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TRANSMITTAL FORM (to be used for all correspondence after initial filing)		Filing Date		ry 29, 20	<u> </u>
		First Named Inventor	Galley		
		Art Unit	1614		
		Examiner Name			
			<u> </u>	Paul V	
Total Number of Pages in This Submission		Attorney Docket Number	21453	} 	
	ENCI	OSURES (Check all that	apply)		
Fee Transmittal Form  Fee Attached  Amendment/Reply  After Final  Affidavits/declaration(s)  Extension of Time Request  Express Abandonment Request  Information Disclosure Statement  Certified Copy of Priority Document(s)  Response to Missing Parts/ Incomplete Application  Response to Missing Parts under 37 CFR 1.52 or 1.53		Drawing(s)  Licensing-related Papers  Petition  Petition to Convert to a  Provisional Application  Power of Attorney, Revocation  Change of Correspondence Addre  Ferminal Disclaimer  Request for Refund  CD, Number of CD(s)	ss	to Group Appeal Com of Appeals a Appeal Com (Appeal Notice Proprietary Status Lette	r sure(s) (please
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## PATENT APPLICATION



## THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application

Inventors:

Galley, et al.

Group: 1614

Serial No. 10/767,784, filed January 29, 2004

Examiner: Ward, Paul V.

(Ref. No. 21453)

For:

MALONAMIDE DERIVATIVES

## RESPONSE TO RESTRICTION REQUIREMENT

Nutley, New Jersey 07110 Date: June 5, 2006

Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

This election is filed in response to the Restriction Requirement issued May 5, 2006, in connection with the above-identified patent application. A response to this Restriction Requirement is due June 5, 2006. Applicants respectfully request consideration of the following remarks.

Claims 1 to 24 are pending. The claims have been restricted to the following forty-two groups.

- I. The compounds according to claim 1 of formula IA, wherein C is phenyl.
- II. The compounds according to claim 1 of formula IA, wherein C is pyridinyl.
- III. The compounds according to claim 1 of formula IA, wherein C is furanyl or tetrahydrofuranyl.

Serial No. 10/767,784 Filed: January 29, 2004

IV.	The compounds according to claim 1 of formula IA, wherein C is benzo[b]thiophenyl.						
V.	The compounds according to claim 1 of formula IA, wherein C is tetrahydronaphthyl.						
VI.	The compounds according to claim 1 of formula IA, wherein C is indanyl.						
VII.	The compounds according to claim 1 of formula IA, wherein C is 2,2,dimethyl-[1,3] dioxolanyl.						
VIII.	The compounds according to claim 1 of formula IB, wherein C is phenyl.						
IX.	The compounds according to claim 1 of formula IB, wherein C is pyridinyl.						
X.	The compounds according to claim 1 of formula IB, wherein C is furanyl or tetrahydrofuranyl.						
XI.	The compounds according to claim 1 of formula IB, wherein C is benzo[b]thiophenyl.						
XII.	The compounds according to claim 1 of formula IB, wherein C is tetrahydronaphthyl.						
XIII.	The compounds according to claim 1 of formula IB, wherein C is indanyl.						
XIV.	The compounds according to claim 1 of formula IB, wherein C is 2,2,dimethyl-[1,3]dioxolanyl.						
XV.	The process for preparing a compound formula IA according to claim 22, wherein C is phenyl.						
XVI.	The process for preparing a compound formula IA according to claim 22, wherein C is pyridinyl.						
XVII.	The process for preparing a compound formula IA according to claim 22, wherein C is furanyl or tetrahydrofuranyl						
XVIII.	The process for preparing a compound formula IA according to claim 22, wherein C is benzo[b]thiophenyl.						
XIX.	The process for preparing a compound formula IA according to claim 22, wherein C is tetrahydronaphthyl.						
XX.	The process for preparing a compound formula IA according to claim 22, wherein C						

is indanyl.

Serial No. 10/767,784 Filed: January 29, 2004

- XXI. The process for preparing a compound formula IA according to claim 22, wherein C 2,2, dimethyl-[1,3] dioxolanyl.
- XXII. The process for preparing a compound formula IB according to claim 23, wherein C is phenyl.
- XXIII. The process for preparing a compound formula IB according to claim 23, wherein C is pyridinyl.
- XXIV. The process for preparing a compound formula IB according to claim 23, wherein C is furanyl or tetrahydrofuranyl.
- XXV. The process for preparing a compound formula IB according to claim 23, wherein C is benzo[b]thiophenyl.
- XXVI. The process for preparing a compound formula IB according to claim 23, wherein C is tetrahydronaphthyl.
- XXVII. The process for preparing a compound formula IB according to claim 23, wherein C is indanyl.
- XXVIII. The process for preparing a compound formula IB according to claim 23, wherein C is 2,2, dimethyl-[1,3]dioxolanyl.
- XXIX. The method of treating according to claims 20-21 of formula IA, wherein C is phenyl.
- XXX. The method of treating according to claims 20-21 of formula IA, wherein C is pyridinyl.
- XXXI. The method of treating according to claims 20-21 of formula IA, wherein C is furanyl or tetrahydrofuranyl.
- XXXII. The method of treating according to claims 20-21 of formula IA, wherein C is benzo[b]thiophenyl.
- XXXIII. The method of treating according to claims 20-21 of formula IA, wherein C is tetrahydronaphthyl.
- XXXIV. The method of treating according to claims 20-21 of formula IA, wherein C is indanyl.
- XXXV. The method of treating according to claims 20-21 of formula IA, wherein C is 2,2, dimethyl-[1,3] dioxolanyl.
- XXXVI. The method of treating according to claims 20-21 of formula IB, wherein C is phenyl.

Serial No. 10/767,784 Filed: January 29, 2004

XXXVII. The method of treating according to claims 20-21 of formula IB, wherein C is

pyridinyl.

XXXVIII. The method of treating according to claims 20-21 of formula IB, wherein C is

furanyl or tetrahydrofuranyl.

XXXIX. The method of treating according to claims 20-21 of formula IB, wherein C is

benzo[b]thiophenyl.

XL. The method of treating according to clams 20-21 of formula IB, wherein C is

tetrahydronaphthyl.

XLI. The method of treating according to claims 20-21 of formula IB, wherein C is

indanyl.

XLII. The method of treating according to claims 20-21 of formula IB, wherein C is 2,2,

dimethyl-[1,3] dioxolanyl.

Applicants hereby elect with traverse Group I, claims 1 to 6, 9 to 14, and 17 to 19, directed to compounds of formula IA wherein C is phenyl and compositions containing them, and the species N-(5-Benzoyl-1-methyl-2-oxo-2,3,4,5-tetrahydro-1H-benzo[b][1,4]diazepin-3-yl)-N'-(3,5-difluoro-benzyl)-2-methyl-malonamide (Example 14), having the following structure:

Claims 1 to 3, 11, 12, and 17 to 19 read on

the elected species.

The requirements are traversed for the following reasons. Upon an election of species, M.P.E.P. § 803.02 states that if no prior art is found that anticipates or renders obvious the elected species, search and examination of the claims will be extended to the extent necessary to

Serial No. 10/767,784

Filed: January 29, 2004

determine patentability of the generic claim. Thus, Applicants understand that if no art is found

which anticipates or renders obvious the elected species, search and examination will be

expanded to the extent necessary to determine patentability of the generic claim or at least to the

extent necessary to determine patentability of Group I.

Further, method claims 20 to 24 have been restricted from the product claims as Groups

XV to XLII. In accordance with M.P.E.P. § 821.04, Applicants understand that upon the finding of

an allowable compound claim, those claims of Groups XV to XLII having all of the limitations of

the allowable product claim will be rejoined and examined on the merits.

The foregoing amendment is fully responsive to the Restriction Requirement issued May 5,

2006. Early and favorable consideration is earnestly solicited.

No additional fees are believed due. However, the Director is hereby authorized to charge

any deficit, or credit any overpayment, to Deposit Account No. 08-2525.

If the Examiner believes there are other issues that can be resolved by telephone interview, or

that there are any informalities remaining in the application which may be corrected by Examiner's

Amendment, a telephone call to the undersigned attorney is respectfully solicited.

Respectfully submitted

Kimberly J. Prior

Attorney for Applicant(s)

(Reg. No. 41,483) 340 Kingsland Street

Nutley, New Jersey 07110

Telephone: (973) 235-6208

Telefax: (973) 235-2363

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